## What we claim is:

## 1. A compound of matter having the formula:

## wherein $R^1$ is $-CH_2OH$ , $-C(=O)NR^5R^6$ ;

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R<sup>2</sup> is selected from the group consisting of hydrogen, C<sub>1-15</sub> alkyl, C<sub>2-15</sub> alkenyl, C<sub>2-15</sub> alkynyl, heterocyclyl, aryl, and heteroaryl, wherein the alkyl, alkenyl, alkynyl, aryl, heterocyclyl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from the group consisting of halo, NO<sub>2</sub>, heterocyclyl, aryl, heteroaryl, CF<sub>3</sub>, CN, OR<sup>20</sup>, SR<sup>20</sup>, N(R<sup>20</sup>)<sub>2</sub>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>)<sub>2</sub>, SO<sub>2</sub>NR<sup>20</sup>COR<sup>22</sup>, SO<sub>2</sub>NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, N(R<sup>20</sup>)<sub>2</sub>, N(R<sup>20</sup>)<sub>2</sub> NR<sup>20</sup>COR<sup>22</sup>, NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, CON(R<sup>20</sup>)<sub>2</sub>, CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, OCONR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, OCON(R<sup>20</sup>)<sub>2</sub>, COON(R<sup>20</sup>)<sub>2</sub>, and OCON(R<sup>20</sup>)<sub>2</sub> and wherein each optional heteroaryl, aryl, and heterocyclyl substituent is optionally substituted with halo, NO<sub>2</sub>, alkyl, CF<sub>3</sub>, amino, mono- or di- alkylamino, alkyl or aryl or heteroaryl amide, NCOR<sup>22</sup>, NR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, COR<sup>20</sup>, CO<sub>2</sub>R<sup>20</sup>, CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, OC(O)R<sup>20</sup>, OC(O)N(R<sup>20</sup>)<sub>2</sub>, SR<sup>20</sup>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>)<sub>2</sub>, CN, and OR<sup>20</sup>;

 $R^3$ ,  $R^4$  are each individually selected from the group consisting of hydrogen,  $C_{1-15}$  alkyl,  $C_{2-15}$  alkenyl,  $C_{2-15}$  alkynyl, heterocyclyl, aryl, and heteroaryl, halo,  $NO_2$ ,  $CF_3$ , CN,  $OR^{20}$ ,  $SR^{20}$ ,  $N(R^{20})_2$ ,  $S(O)R^{22}$ ,  $SO_2R^{22}$ ,  $SO_2N(R^{20})_2$ ,  $SO_2NR^{20}COR^{22}$ ,  $SO_2NR^{20}CO_2R^{22}$ ,  $SO_2NR^{20}CON(R^{20})_2$ ,  $N(R^{20})_2$ , wherein the alkyl, alkenyl, alkynyl, aryl, heterocyclyl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from the group consisting of halo,  $NO_2$ , heterocyclyl, aryl, heteroaryl,  $N(R^{20})_2$ ,  $N(R^{20})_2$ ,

SO<sub>2</sub>NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, N(R<sup>20</sup>)<sub>2</sub> NR<sup>20</sup>COR<sup>22</sup>, NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, CONR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, NR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, OCONR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, OC(O)R<sup>20</sup>, C(O)OCH<sub>2</sub>OC(O)R<sup>20</sup>, and OCON(R<sup>20</sup>)<sub>2</sub> and wherein each optional heteroaryl, aryl, and heterocyclyl substituent is optionally substituted with halo, NO<sub>2</sub>, alkyl, CF<sub>3</sub>, amino, mono- or di- alkylamino, alkyl or aryl or heteroaryl amide, NCOR<sup>22</sup>, NR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, COR<sup>20</sup>, CO<sub>2</sub>R<sup>20</sup>, CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, OC(O)R<sup>20</sup>, OC(O)N(R<sup>20</sup>)<sub>2</sub>, SR<sup>20</sup>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>)<sub>2</sub>, CN, and OR<sup>20</sup>;

R<sup>5</sup> and R<sup>6</sup> are each individually selected from H, C<sub>1-15</sub> alkyl with from 1 to 2 substituents independently selected from the group consisting of halo, NO<sub>2</sub>, heterocyclyl, aryl, heteroaryl, CF<sub>3</sub>, CN, OR<sup>20</sup>, SR<sup>20</sup>, N(R<sup>20</sup>)<sub>2</sub>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>)<sub>2</sub>, SO<sub>2</sub>NR<sup>20</sup>COR<sup>22</sup>, SO<sub>2</sub>NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, N(R<sup>20</sup>)<sub>2</sub>, N(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, CON(R<sup>20</sup>)<sub>2</sub>, CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>NR<sup>20</sup>CO<sub>2</sub>R<sup>22</sup>, OCONR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, OC(O)R<sup>20</sup>, C(O)OCH<sub>2</sub>OC(O)R<sup>20</sup>, and OCON(R<sup>20</sup>)<sub>2</sub>, and wherein each optional heteroaryl, aryl, and heterocyclyl substituent is optionally substituted with halo, NO<sub>2</sub>, alkyl, CF<sub>3</sub>, amino, mono- or di- alkylamino, alkyl or aryl or heteroaryl amide, NCOR<sup>22</sup>, NR<sup>20</sup>SO<sub>2</sub>R<sup>22</sup>, COR<sup>20</sup>, CO<sub>2</sub>R<sup>20</sup>, CON(R<sup>20</sup>)<sub>2</sub>, NR<sup>20</sup>CON(R<sup>20</sup>)<sub>2</sub>, OC(O)R<sup>20</sup>, OC(O)N(R<sup>20</sup>)<sub>2</sub>, SR<sup>20</sup>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>), CN, and OR<sup>20</sup>;

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 $R^{20}$  is selected from the group consisting of H,  $C_{1-15}$  alkyl,  $C_{2-15}$  alkenyl,  $C_{2-15}$  alkynyl, heterocyclyl, aryl, and heteroaryl, wherein the alkyl, alkenyl, alkynyl, heterocyclyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from halo, alkyl, mono- or dialkylamino, alkyl or aryl or heteroaryl amide, CN, O- $C_1$  alkyl, CF<sub>3</sub>, aryl, and heteroaryl; and

 $R^{22}$  is a member selected from the group consisting of  $C_{1-15}$  alkyl,  $C_{2-15}$  alkenyl,  $C_{2-15}$  alkynyl, heterocyclyl, aryl, and heteroaryl, wherein the alkyl, alkenyl, alkynyl, heterocyclyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from halo, alkyl, mono- or dialkylamino, alkyl or aryl or heteroaryl amide, CN, O- $C_{1-6}$  alkyl, CF<sub>3</sub>, and heteroaryl wherein, when  $R^1$  is CH<sub>2</sub>OH, and  $R^3$  is H and  $R^4$  is H, and the pyrazole ring is attached through  $C^4$ , then  $R^2$  is not H.

2. The compound of claim 1 wherein R<sup>2</sup> is selected from the group consisting of hydrogen, C<sub>1-15</sub> alkyl, C<sub>2-15</sub> alkynyl, heterocyclyl, aryl, and heteroaryl, wherein the alkyl, alkynyl, aryl, heterocyclyl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from the group consisting of halo, NO<sub>2</sub>, heterocyclyl, aryl, heteroaryl, CF<sub>3</sub>, CN, OR<sup>20</sup>, SR<sup>20</sup>, N(R<sup>20</sup>)<sub>2</sub>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>)<sub>2</sub>, COR<sup>20</sup>, CO<sub>2</sub>R<sup>20</sup>, CON(R<sup>20</sup>)<sub>2</sub>, and wherein each optional heteroaryl, aryl, and heterocyclyl substituent is

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optionally substituted with halo, alkyl, CF, CN, and OR<sup>20</sup>;

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, C<sub>1.15</sub> alkyl, C<sub>2-15</sub> alkynyl, heterocyclyl, aryl, heteroaryl, halo, NO<sub>2</sub>, CF<sub>3</sub>, CN, OR<sup>20</sup>, SR<sup>20</sup>, N(R<sup>20</sup>)<sub>2</sub>, S(O)R<sup>22</sup>, SO<sub>2</sub>R<sup>22</sup>, SO<sub>2</sub>N(R<sup>20</sup>)<sub>2</sub>, COR<sup>20</sup>, CO<sub>2</sub>R<sup>20</sup>, CON(R<sup>20</sup>)<sub>2</sub>, wherein the alkyl, alkynyl, aryl, heterocyclyl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from the group consisting of halo, NO<sub>2</sub>, heterocyclyl, aryl, heteroaryl,  $CF_3$ , CN,  $OR^{20}$ ,  $SR^{20}$ ,  $N(R^{20})_2$ ,  $S(O)R^{22}$ ,  $SO_2R^{22}$ ,  $SO_2N(R^{20})_2$ ,  $COR^{20}$ ,  $CO_2R^{20}$ , CON(R<sup>20</sup>), and wherein each optional heteroaryl, aryl, and heterocyclyl substituent is optionally substituted with halo, alkyl, CF, CN, and OR<sup>20</sup>;

R<sup>5</sup> and R<sup>6</sup> are each individually selected from H, and C<sub>1-15</sub> alkyl having from 1 to 2 substituents independently selected from the group consisting of aryl, heteroaryl, CF<sub>3</sub>, OR<sup>20</sup>, and wherein each optional heteroaryl, and aryl substituent is further optionally substituted with halo, alkyl, and CF<sub>3</sub>;

R<sup>20</sup> is a member selected from the group consisting of H, C<sub>1-6</sub> alkyl, aryl, and heteroaryl; and

 $R^{22}$  is a member selected from the group consisting of  $C_{1.6}$  alkyl, aryl, and heteroaryl, wherein the alkyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from halo, alkyl, CN, O-C<sub>1-6</sub> alkyl, and CF<sub>3</sub>.

3. The compound of claim 1 wherein R<sup>2</sup> is selected from the group consisting of hydrogen, C<sub>1-15</sub> alkyl, C<sub>2-15</sub> aryl, and heteroaryl, wherein the alkyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from the group consisting of halo, aryl, heteroaryl, CF<sub>3</sub>, CN, OR<sup>20</sup>, SR<sup>20</sup>, S(O)R<sup>22</sup>, CO<sub>2</sub>R<sup>20</sup>, CON(R<sup>20</sup>)<sub>2</sub>, and wherein each optional heteroaryl, and aryl substituent is optionally substituted with halo, alkyl, CF, CN, and OR<sup>20</sup>;

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, C<sub>1,15</sub> alkyl,  $C_{2-15}$  aryl, heteroaryl, halo,  $CF_3$ , CN,  $OR^{20}$ ,  $SR^{20}$ ,  $S(O)R^{22}$ ,  $CO_2R^{20}$ , and  $CON(R^{20})_2$ , wherein the alkyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 3 substituents independently selected from the group consisting of halo, aryl, heteroaryl, CF<sub>3</sub>, CN, OR<sup>20</sup>, SR<sup>20</sup>, S(O)R<sup>22</sup>, CO<sub>2</sub>R<sup>20</sup>, and CON(R<sup>20</sup>)<sub>2</sub>, and wherein each optional heteroaryl, and aryl substituent is optionally substituted with halo, alkyl, CF, CN, and OR<sup>20</sup>;

R<sup>5</sup> and R<sup>6</sup> are each individually selected from H, and C<sub>1-15</sub> alkyl having from 1 to 2 substituents selected from CF,;

R<sup>20</sup> is selected from H, and C<sub>1-6</sub>; and  $R^{22}$  is  $C_{1.6}$  alkyl.

4. The compound of claim 1 wherein  $R^2$  is independently selected from the group consisting of hydrogen,  $C_{1-15}$  alkyl,  $C_{2-15}$  aryl, and heteroaryl, wherein the alkyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, aryl, heteroaryl,  $CF_3$ , CN,  $OR^{20}$ ,  $CO_2R^{20}$ , and  $CON(R^{20})_2$ , and wherein each optional heteroaryl, and aryl substituent is optionally substituted with halo, alkyl,  $CF_3$  and CN;

 $R^3$  and  $R^4$  are each individually selected from the group consisting of hydrogen,  $C_{1.15}$  alkyl,  $C_{2.15}$  aryl, heteroaryl, halo,  $CF_3$ , CN,  $OR^{20}$ ,  $CO_2R^{20}$ , and  $CON(R^{20})_2$ , wherein the alkyl, aryl, and heteroaryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, aryl, heteroaryl,  $CF_3$ , CN,  $OR^{20}$ ,  $CO_2R^{20}$ , and  $CON(R^{20})_2$ , and wherein each optional heteroaryl, and aryl substituent is optionally substituted with halo, alkyl,  $CF_3$  or CN;

 $R^{5}$  and  $R^{6}$  are each individually selected from H, and  $C_{1-15}$  alkyl;

R<sup>20</sup> is selected from H, and C<sub>1-6</sub>; and

 $R^{22}$  is  $C_{1-6}$  alkyl.

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5. The compound of claim 1 wherein R<sup>2</sup> is independently selected from the group consisting of hydrogen, C<sub>1-15</sub> alkyl and aryl, wherein the alkyl and aryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, OR<sup>20</sup>, aryl, CF<sub>3</sub>, CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> or CN;

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, C<sub>1-15</sub> alkyl, aryl, halo, CF<sub>3</sub>, and CN, wherein the alkyl, and aryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, aryl, CF<sub>3</sub>, CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> or CN;

 $R^5$  and  $R^6$  are each individually selected from H, and  $C_{1-15}$  alkyl; and  $R^{20}$  is selected from H, and  $C_{1-6}$ .

6. The compound of claims 1 or 2 or 3 or 4 or 5 having the following formula wherein the point of attachment of the pyrazole ring is C-4.

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7. The compound of claims 1 or 2 or 3 or 4 or 5 having the following formula wherein the point of attachment of the pyrazole ring is C-3.

8. The compound of claims 1 or 2 or 3 or 4 or 5 having the following formula 5 wherein the point of attachment of the pyrazole ring is C-5.

- 9. The compound of claims 6 or 7 or 8 wherein  $R^1 = CH_2OH$ ;
- 10. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is independently selected from the group consisting of hydrogen, C<sub>1-10</sub> alkyl and aryl, wherein the alkyl and aryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, OR<sup>20</sup>, aryl, CF<sub>3</sub>, and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN;

R<sup>3</sup> and R<sup>4</sup> are each individually- selected from the group consisting of hydrogen, C<sub>1-15</sub> alkyl, aryl, halo, CF<sub>3</sub>, and CN, wherein the alkyl, and aryl substituents are optionally substituted with a substituent independently selected from the group consisting of halo, CF<sub>3</sub>, and CN; and

R<sup>20</sup> is selected from H, and C<sub>1-6</sub> alkyl;

11. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is independently selected from the group consisting of hydrogen, C<sub>1.8</sub> alkyl and aryl, wherein the alkyl and aryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, OR<sup>20</sup>, aryl, CF<sub>3</sub>, and

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CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN;

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, C<sub>1.3</sub> alkyl, aryl, halo, CF<sub>3</sub>, CN; and

R<sup>20</sup> is selected from H, and C<sub>1-6</sub> alkyl.

12. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is independently selected from the group consisting of hydrogen, C<sub>1.8</sub> alkyl and aryl, wherein the alkyl and aryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, OR<sup>20</sup>, aryl, CF<sub>3</sub>, and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN;

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, methyl, and halo; and

R<sup>20</sup> is selected from H, and C<sub>1-6</sub>;

13. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is independently selected from the group consisting of hydrogen, C<sub>1-8</sub> alkyl and aryl, wherein the alkyl and aryl substituents are optionally substituted with 1 substituent selected from the group consisting of halo, aryl, CF<sub>3</sub>, and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, and methyl.

14. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1-8</sub> alkyl that is optionally substituted with 1 substituent selected from the group consisting of aryl, CF<sub>3</sub>, and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, and methyl.

15. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1.8</sub> alkyl that is optionally substituted with one aryl substituent that is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

R<sup>3</sup> and R<sup>4</sup> are each hydrogen.

16. The compound of claim 6 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

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 $R^2$  is selected from the group consisting of hydrogen, and  $C_{1-6}$  alkyl that is optionally substituted with aryl that is optionally substituted with alkyl; and

R³ and R⁴ are each hydrogen.

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17. The compound of claim 7 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

 $R^2$  is selected from the group consisting of hydrogen, and  $C_{1-8}$  alkyl that is optionally substituted with 1 substituent selected from the group consisting of aryl,  $CF_3$ , and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl,  $CF_3$  and CN; and

R<sup>3</sup> and R<sup>4</sup> are each individually selected from hydrogen, and methyl.

18. The compound of claim 7 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1.8</sub> alkyl that is optionally substituted with 1 substituent selected from the group consisting of aryl, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

R³ and R⁴ are each hydrogen.

19. The compound of claim 7 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

 $R^2$  is selected from the group consisting of hydrogen, and  $C_{1-6}$  alkyl that is optionally substituted with aryl that is optionally substituted with alkyl; and

R<sup>3</sup> and R<sup>4</sup> are each hydrogen.

20. The compound of claim 8 wherein R' is -CH<sub>2</sub>OH;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1-6</sub> alkyl;

R<sup>3</sup> is selected from the group consisting of hydrogen, C<sub>1-6</sub> alkyl, and aryl, wherein the alkyl, and aryl substituents are optionally substituted with from 1 to 2 substituents independently selected from the group consisting of halo, aryl, CF<sub>3</sub>, and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

 $R^4$  is selected from the group consisting of hydrogen and  $C_{1.6}$  alkyl.

21. The compound of claim 8 wherein R<sup>1</sup> is -CH<sub>2</sub>OH;

R<sup>2</sup> is selected from the group consisting of hydrogen, and methyl;

 $R^3$  and  $R^4$  are each independently selected from the group consisting of hydrogen, and  $C_{1-6}$  alkyl, that is optionally substituted with aryl that is optionally substituted with alkyl; and

R<sup>4</sup> is selected from hydrogen and methyl.

22. The compound of claim 6 wherein R<sup>1</sup> is -CONHEt;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1.8</sub> alkyl that is optionally substituted with 1 substituent selected from the group consisting of aryl, CF<sub>3</sub>, and CN, and wherein each optional aryl substituent is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN;

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and

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R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, and methyl.

23. The compound of claim 6 wherein R<sup>1</sup> is -CONHEt;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1.8</sub> alkyl that is optionally substituted with 1 aryl substitutent of aryl, that is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

R<sup>3</sup> and R<sup>4</sup> are each hydrogen.

24. The compound of claim 6 wherein R<sup>1</sup> is -CONHEt;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1-6</sub> alkyl that is optionally substituted with aryl that is optionally substituted with alkyl; and

R³ and R⁴ are hydrogen.

25. The compound of claim 7 wherein R<sup>1</sup> is -CONHEt;

R<sup>2</sup> is selected from the group consisting of hydrogen, and C<sub>1.8</sub> that is optionally substituted with 1 aryl substituent that is optionally substituted with halo, alkyl, CF<sub>3</sub> and CN; and

R³ and R⁴ are hydrogen.

26. The compound of claim 7 wherein R<sup>1</sup> is -CONHEt;

 $R^2$  is independently selected from the group consisting of hydrogen, and  $C_{1-6}$  alkyl that is optionally substituted with aryl that is optionally substituted with alkyl; and

R<sup>3</sup> and R<sup>4</sup> are each hydrogen.

27. The compound of claim 8 wherein R<sup>1</sup> is -CONHEt; and

R<sup>2</sup> is selected from hydrogen, and methyl;

R<sup>3</sup> and R<sup>4</sup> are each individually selected from the group consisting of hydrogen, and C<sub>1-6</sub> alkyl, wherein the alkyl, is optionally substituted with aryl that is optionally substituted with alkyl; and

R<sup>4</sup> is selected from hydrogen and methyl.

- 28. The compound of claim 1 wherein the compound is selected from (4S,2R,3R,5R)-2-{6-amino-2-[1-benzylpyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol,
- 30 (4S,2R,3R,5R)-2-[6-amino-2-(1-pentylpyrazol-4-yl)purin-9yl]-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-[6-amino-2-(1-methylpyrazol-4-yl)purin-9-yl]-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-(methylethyl)pyrazol-4-yl)pyrazol-4-yl)pyrazol-4-

yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-(3phenylpropyl)pyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-(4-t-butylbenzyl)pyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-(6-amino-2-pyrazol-4-ylpurin-9-yl)-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-pent-4-enylpyrazol-4-v1]purin-9-v1}-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-decylpyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol. (4S,2R,3R,5R)-2-{6-amino-2-[1-(cyclohexylmethyl)pyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-(2-phenylethyl)pyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol, 10 (4S,2R,3R,5R)-2-{6-amino-2-[1-(3cyclohexylpropyl)pyrazol-4-yl]purin-9-yl}-5-(hydroxymethyl)oxolane-3,4-diol, (4S,2R,3R,5R)-2-{6-amino-2-[1-(2-cyclohexylethyl)pyrazol-4-yl]purin-9-yl}-5-

- 29. A method for stimulating coronary vasodilatation in a mammal by administering to the mammal a therapeutically effective amount of a compound of claim 1 that is sufficient to stress the heart and induce a coronary steal situation for the purposes of imaging the heart.
  - 30. The method of claim 25 wherein the therapeutically effective amount ranges from about 0.01 to about 100 mg/kg weight of the mammal.
    - 31. The method of claim 29 wherein the mammal is a human.

(hydroxymethyl)oxolane-3,4-diol, and mixtures thereof.

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- 32. A pharmaceutical composition of matter comprising the compound of claim 1 and one or more pharmaceutical excipients.
- 33. The pharmaceutical composition of matter of claim 32 wherein the pharmaceutical composition is in the form of a solution.
- 34. The pharmaceutical composition of matter of claim 32 wherein the composition is useful as an anti-inflammatory, in adjunctive therapy with angioplasty, as a platelet aggregation inhibitor, and as an inhibitor of platelet and neutrophil activation.